**AUSTRALIAN** \*Clinicalabs LABORATORY 3427-18703 Referred: 21/07/21

UNIT 104/14 LEXINGTON D Collected: 22/07/21 12:40
BELLA VISTA Tested: 22/07/21
Tel:1300 134 111 Printed: 07/09/21 18:14

Batch: 12082 1

DR RACHAEL RODGERS MR EMMETT, MATTHEW

6 WATKINS ROAD |FERTILITY SPECIALIST SYDNEY/B UR Ref : |SE 1204, LVL 12, 135-137 MACQ AVALON 2107

DOB: 20/03/1991 (30 Y) Lab No:21-51555514-I SYDNEY

\_\_\_\_Sex :Male \_

Ph: 0414194286 | Sex :Male | Clinical Notes: fertility assessment

## TEST NAME: KARYOTYPE STUDIES

This test was performed by: Children's Hospital at Westmead, Level 2, Diagnostic building Cnr Hawkesbury & Hamsworth Rd. WESTMEAD NSW 2145 Ph: (02)9845 3276 Fax: (02) 9845 3332

Test was referred on 23/07/21

Our records show a hard copy of this report has been dispatched to the surgery

KAR-R, HCR-R, TRP-R, IS-R, HEP-R, FBE-R, HIV-R, HBE-R CC Drs: PATIENT COPY:.

MR EMMETT, MATTHEW KARYOTYPE STUDIES

Page: 1 of 1 All Tests Complete





attn. Dr. RODGERS.RACHAEL Australian Clinical Laboratories NSW Unit 108 14 Lexington Drive BELLA VISTA NSW 2153

Corner Hawkesbury Boad & Hainsworth Street Locked Bag 4001 Westmead Sydney NSW 2145 Australia Telephone (02)9845-3237 Facsimile (02)9845-3238

MRN: 50195349 EMMETT, Matthew

6 Watkins Rd AVALON BEACH INSW 2107 DOB: 20/03/1991 Sex.Male Client MRN: NA

Accession No: CG-21-06789

## Cytogenetics

## Cytogenetics Final Report

Accession #: CG-21-06789 Collection date: 22/07/2021 12:40 Received date: 23/07/2021 15:12

Specimen type: Blood

External Reference: 21-51555514 Clinical indication: Fertility assessment

Test Requested: Cytogenetic analysis referred for "Karyotype"

Preparation: PHA stimulated lymphocyte culture.

Resolution: 400-550bphs Panding: G Banding.

Total cells: 5 Cells counted: 0 Cells scored: 0 Cells analysed: 5

Karyotype Result: 46,XY

## Conclusion:

A 46,XY male karyotype with no cytogenetic abnormality.

Electronically signed by Praveen Sharma, MSc. MHGSA (Cytogenetics) √erified 26.08.21

Limitation: This karyotype test has been performed to exclude a structural chromosome rearrangement at the above stated G-band resolution Smaller chromosome abnormalities <5-10Mb genomic content may go undetected. It has limited detection for mosaicism and does not detect. eniparental disorny. DNA sequence variants or triplet repeat disorders. This result is reported and verified by Mr.P. Sharma MSc. MHGSA (Serior Scientist, Cytogenetics Dept, CHW), under authorisation of Mr D Wright (Head of Dept, Cytogenetics, CHW),

Air Dale Wright

Head of Department

Enquines +61 2 98453237

Mr Luke St Heaps

Deputy Head of Department

Dr Anja Ravine

Genetic Pathologist

Provider Number 035083EL





Page 1 of 1

NATA/RCPA Accredited Laboratory: 2254

Accredited for compliance with NPAAC Standards and ISO 15189

Printed: 26/08/2021 14:10

Report ID: 41133360

All data contained in this report is performed within the SCHN eMR and constitute part of the medical record in accordance with NSW Meanth perfor